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FILE 'USPATFULL' ENTERED AT 15:40:11 ON 09 JUN 2003  
CA INDEXING COPYRIGHT (C) 2003 AMERICAN CHEMICAL SOCIETY (ACS)

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=> s opa1  
L1 187 OPA1

=> s l1 and glia#  
L2 3 L1 AND GLIA#

=> s l1 and neur?  
L3 43 L1 AND NEUR?

=> duplicate remove  
ENTER L# LIST OR (END):12  
DUPLICATE PREFERENCE IS 'USPATFULL, PCTFULL'  
KEEP DUPLICATES FROM MORE THAN ONE FILE? Y/(N):n  
PROCESSING COMPLETED FOR L2  
L4 3 DUPLICATE REMOVE L2 (0 DUPLICATES REMOVED)

=> duplicate remove  
ENTER L# LIST OR (END):13  
DUPLICATE PREFERENCE IS 'MEDLINE, BIOSIS, USPATFULL, PCTFULL'  
KEEP DUPLICATES FROM MORE THAN ONE FILE? Y/(N):n  
PROCESSING COMPLETED FOR L3  
L5 32 DUPLICATE REMOVE L3 (11 DUPLICATES REMOVED)

=> d 1-32

L5 ANSWER 1 OF 32 MEDLINE  
AN 2002725528 MEDLINE  
DN 22375944 PubMed ID: 12488251  
TI The OPA1 gene and optic neuropathy.  
CM Comment on: Br J Ophthalmol. 2003 Jan;87(1):48-53  
AU Alward W L M  
SO BRITISH JOURNAL OF OPHTHALMOLOGY, (2003 Jan) 87 (1) 2-3.  
Journal code: 0421041. ISSN: 0007-1161.  
CY England: United Kingdom  
DT Commentary  
Editorial  
LA English  
FS Priority Journals  
EM 200302  
ED Entered STN: 20021219  
Last Updated on STN: 20030221  
Entered Medline: 20030220

L5 ANSWER 2 OF 32 USPATFULL  
AN 2003:120036 USPATFULL  
TI Cellular kinases involved in Cytomegalovirus infection and their inhibition  
IN Schubart, Daniel, Weil am Rhein, GERMANY, FEDERAL REPUBLIC OF  
Habenberger, Peter, Munchen, GERMANY, FEDERAL REPUBLIC OF  
Stein-Gerlach, Matthias, Munchen, GERMANY, FEDERAL REPUBLIC OF

FS APPLICATION  
LN.CNT 2300  
INCL INCLM: 435/905,000

NCL INCLS: 424/146.100; 514/.000; 435/007.100; 435/069.100  
NCLM: 435/005.000  
NCLS: 424/146.100; 514/001.000; 435/007.100; 435/069.100  
IC [7]  
ICM: A61K039-395  
ICS: C12Q001-70; A61K031-00; A01N061-00; G01N033-53; C12P021-06  
CAS INDEXING IS AVAILABLE FOR THIS PATENT.

L5 ANSWER 3 OF 32 USPATFULL  
AN 2003:44371 USPATFULL  
TI Combined growth factor-deleted and thymidine kinase-deleted vaccinia virus vector  
IN McCart, J. Andrea, Toronto, CANADA  
Bartlett, David L., Pittsburgh, PA, UNITED STATES  
Moss, Bernard, Bethesda, MD, UNITED STATES  
PI US 2003031681 A1 20030213  
AI US 2001-991721 A1 20011113 (9)  
PRAI WO 2000-US14679 20000526  
US 1999-137126P 19990528 (60)  
DT Utility  
FS APPLICATION  
LN.CNT 2762  
INCL INCLM: 424/186.100  
INCLS: 435/456.000; 435/235.100  
NCL NCLM: 424/186.100  
NCLS: 435/456.000; 435/235.100  
IC [7]  
ICM: A61K039-12  
ICS: C12N015-863; C12N007-00  
CAS INDEXING IS AVAILABLE FOR THIS PATENT.

L5 ANSWER 4 OF 32 USPATFULL  
AN 2003:3547 USPATFULL  
TI Trans-viral vector mediated gene transfer to the retina  
IN Wakefield, John, Birmingham, AL, UNITED STATES  
Bennett, Jean, Bryn Mawr, PA, UNITED STATES  
PA Tranzyme, Inc. (U.S. corporation)  
PI US 2003003582 A1 20030102  
AI US 2002-140227 A1 20020507 (10)  
PRAI US 2001-340116P 20011102 (60)  
US 2001-289459P 20010508 (60)  
DT Utility  
FS APPLICATION  
LN.CNT 2316  
INCL INCLM: 435/456.000  
INCLS: 424/093.200  
NCL NCLM: 435/456.000  
NCLS: 424/093.200  
IC [7]  
ICM: A61K048-00  
ICS: C12N015-867  
CAS INDEXING IS AVAILABLE FOR THIS PATENT.

L5 ANSWER 5 OF 32 MEDLINE DUPLICATE 1  
AN 2003114334 MEDLINE  
DN 22499677 PubMed ID: 12509422  
TI Loss of **OPA1** perturbs the mitochondrial inner membrane structure and integrity, leading to cytochrome c release and apoptosis.  
AU Olichon Aurelien; Baricault Laurent; Gas Nicole; Guillou Emmanuelle;  
Valette Annie; Belenguer Pascale; Lenaers Guy  
CS Laboratoire de Biologie Cellulaire et Moleculaire du Controle de la ..

United States  
DT Journal; Article; (JOURNAL ARTICLE)  
LA English  
FS Priority Journals

OS OMIM-165500  
EM 200304  
ED Entered STN: 20030312  
Last Updated on STN: 20030424  
Entered Medline: 20030423

L5 ANSWER 6 OF 32 MEDLINE DUPLICATE 2  
AN 2002725536 MEDLINE  
DN 22375955 PubMed ID: 12488262  
TI Optic disc morphology of patients with **OPA1** autosomal dominant optic atrophy.  
CM Comment in: Br J Ophthalmol. 2003 Jan;87(1):2-3  
AU Votruba M; Thiselton D; Bhattacharya S S  
CS Department of Molecular Genetics, Institute of Ophthalmology, UCL, Bath Street, London EC1V 9EL, UK.. m\_votruba@altavista.co.uk  
SO BRITISH JOURNAL OF OPHTHALMOLOGY, (2003 Jan) 87 (1) 48-53.  
Journal code: 0421041. ISSN: 0007-1161.  
CY England: United Kingdom  
DT Journal; Article; (JOURNAL ARTICLE)  
LA English  
FS Priority Journals  
EM 200302  
ED Entered STN: 20021219  
Last Updated on STN: 20030221  
Entered Medline: 20030220

L5 ANSWER 7 OF 32 USPATFULL  
AN 2002:314675 USPATFULL  
TI COX 1-interacting proteins and use thereof  
IN Wettstein, Daniel Albert, Salt Lake City, UT, UNITED STATES  
PA Myriad Genetics, Incorporated, Salt Lake City, UT (U.S. corporation)  
PI US 2002177152 A1 20021128  
AI US 2002-100503 A1 20020318 (10)  
PRAI US 2001-277013P 20010319 (60)  
DT Utility  
FS APPLICATION  
LN.CNT 4721  
INCL INCLM: 435/006.000  
INCLS: 435/069.100; 435/189.000; 435/320.100; 435/325.000  
NCL NCLM: 435/006.000  
NCLS: 435/069.100; 435/189.000; 435/320.100; 435/325.000  
IC [7]  
ICM: C12Q001-68  
ICS: C12N009-02  
CAS INDEXING IS AVAILABLE FOR THIS PATENT.

L5 ANSWER 8 OF 32 PCTFULL COPYRIGHT 2003 Univentio  
AN 2002027022 PCTFULL ED 20020701 EW 200214  
TIEN IMPROVEMENTS IN AND RELATING TO TREATMENTS FOR EYE DISEASE  
TIFR AMELIORATIONS RELATIVES AUX TRAITEMENTS DE MALADIE DES YEUX  
IN BHATTACHARYA, Shomi, University College London, Institute of Ophthalmology, 11-43 Bath Street, London EC1V 9EL, GB [GB, GB]; WISSINGER, Bernd, University Eye Hospital, Auf der Morgenstelle 15, 27076 Tubingen, DE [DE, DE]; ALEXANDER, Christiana, University Eye Hospital, Auf der Morgenstelle 15, 27076 Tubingen, DE [DE, DE]; VOTRUBA, Marcela, University College London, Institute of Ophthalmology, 11-43 Bath Street, London EC1V 9EL, GB [GB, GB]  
PA UNIVERSITY COLLEGE LONDON, Institute of Ophthalmology, 11-43 Bath Street, London EC1V 9EL, GB [GB, GB], for all designates States except US;

WISSINGER, Bernd, Universit; Eye Hospital, Auf der Morgenstelle 15, 27076 Tubingen, DE [DE, DE], for US only;  
ALEXANDER, Christiana, University Eye Hospital, Auf der Morgenstelle 15,

27076 Tubingen, DE [DE, , for US only;  
VOTRUBA, Marcela, University College London, Institute of Ophthalmology,  
11-43 Bath Street, London EC1V 9EL, GB [GB, GB], for US only  
AG SILVESTON, Judith, Abel & Imray, 20 Red Lion Street, London WC1R 4PQ, GB  
LAF English  
LA English  
DT Patent  
PI WO 2002027022 A2 20020404  
DS W: AE AG AL AM AT AU AZ BA BB BG BR BY BZ CA CH CN CO CR CU CZ  
DE DK DM DZ EC EE ES FI GB GD GE GH GM HR HU ID IL IN IS JP  
KE KG KP KR KZ LC LK LS LT LU LV MA MD MG MK MN MW MX MZ  
NO NZ PH PL PT RO RU SD SE SG SI SK SL TJ TM TR TT TZ UA UG  
US UZ VN YU ZA ZW  
RW (ARIPO): GH GM KE LS MW MZ SD SL SZ TZ UG ZW  
RW (EAPO): AM AZ BY KG KZ MD RU TJ TM  
RW (EPO): AT BE CH CY DE DK ES FI FR GB GR IE IT LU MC NL PT SE TR  
RW (OAPI): BF BJ CF CG CI CM GA GN GQ GW ML MR NE SN TD TG  
AI WO 2001-GB4284 A 20010926  
PRAI GB 2000-0023555.6 20000926  
ICM C12Q001-68

L5 ANSWER 9 OF 32 PCTFULL COPYRIGHT 2003 Univentio  
AN 2002000878 PCTFULL ED 20020814  
TIEN HUMAN MSP1 MITOCHONDRIAL DYNAMIN, ITS MSP1-X ISOFORMS, AND THEIR  
THERAPEUTIC USE  
TIFR DYNAMINE MITOCHONDRIALE HUMAINE MSP1, SES ISOFORMES MSP1-X, ET LEUR  
UTILISATION EN THERAPEUTIQUE  
IN LENAERS, Guy;  
DUCOMMUN, Bernard;  
HAMEL, Christian;  
DELETTRE, Cecile;  
BELENGUER, Pascale  
PA UNIVERSITE PAUL SABATIER;  
INSTITUT NATIONAL DE LA SANTE ET DE LA RECHERCHE MEDICALE;  
LENAERS, Guy;  
DUCOMMUN, Bernard;  
HAMEL, Christian;  
DELETTRE, Cecile;  
BELENGUER, Pascale  
DT Patent  
PI WO 2002000878 A2 20020103  
DS W: US AT BE CH CY DE DK ES FI FR GB GR IE IT LU MC NL PT SE TR  
AI WO 2001-FR1999 A 20010625  
PRAI FR 2000-00/08140 20000626  
ICM C12N015-12  
ICS C12N015-63; C12N001-19; C12N001-21; C07K014-47; C07K016-18;  
A61K038-17

L5 ANSWER 10 OF 32 MEDLINE DUPLICATE 3  
AN 2002260122 MEDLINE  
DN 21975182 PubMed ID: 11847212  
TI Primary structure of a dynamin-related mouse mitochondrial GTPase and its  
distribution in brain, subcellular localization, and effect on  
mitochondrial morphology.  
AU Misaka Takumi; Miyashita Tomoyuki; Kubo Yoshihiro  
CS Department of Physiology, Tokyo Medical and Dental University, Graduate  
School and Faculty of Medicine, Bunkyo, Tokyo 113-8519, Japan.  
SO JOURNAL OF BIOLOGICAL CHEMISTRY, (2002 May 3) 277 (18) 15834-42.  
CY Journal code: 2985121R. ISSN: 0021-9258.  
PT United States  
Journal; Article; (JOURNAL ARTICLE)

Entered STN: 20020702  
Last Updated on STN: 20030105  
Entered Medline: 20020702

L5 ANSWER 11 OF 32 MEDLINE  
AN 2002680827 IN-PROCESS  
DN 22328899 PubMed ID: 12441838  
TI Is normal tension glaucoma actually an unrecognized hereditary optic neuropathy? New evidence from genetic analysis.  
AU Buono Lawrence M; Foroozan Rod; Sergott Robert C; Savino Peter J  
SO CURRENT OPINION IN OPHTHALMOLOGY, (2002 Dec) 13 (6) 362-70.  
Journal code: 9011108. ISSN: 1040-8738.  
CY United States  
DT Journal; Article; (JOURNAL ARTICLE)  
LA English  
FS IN-PROCESS; NONINDEXED; Priority Journals; Health Technology  
ED Entered STN: 20021121  
Last Updated on STN: 20021213

L5 ANSWER 12 OF 32 MEDLINE DUPLICATE 4  
AN 2002148751 MEDLINE  
DN 21846718 PubMed ID: 11855928  
TI OPA1 (Kjer type) dominant optic atrophy: a novel mitochondrial disease.  
AU Delettre Cecile; Lenaers Guy; Pelloquin Laeticia; Belenguer Pascale; Hamel Christian P  
CS Inserm U.254, Laboratoire de Neurobiologie de l'audition, 71, rue de Navacelles, 34090Montpellier, France.. delettre@montp.inserm.fr  
SO MOLECULAR GENETICS AND METABOLISM, (2002 Feb) 75 (2) 97-107. Ref: 71  
Journal code: 9805456. ISSN: 1096-7192.  
CY United States  
DT Journal; Article; (JOURNAL ARTICLE)  
General Review; (REVIEW)  
(REVIEW, TUTORIAL)  
LA English  
FS Priority Journals  
EM 200205  
ED Entered STN: 20020308  
Last Updated on STN: 20020522  
Entered Medline: 20020521

L5 ANSWER 13 OF 32 BIOSIS COPYRIGHT 2003 BIOLOGICAL ABSTRACTS INC.  
AN 2003:155358 BIOSIS  
DN PREV200300155358  
TI Lack of Phenotypic Differences in Normal Tension Glaucoma Patients With and Without OPA1 Polymorphisms.  
AU Okada, K. (1); Aung, T.; Poinoosawmy, D.; Brice, G.; Child, A. H.; Bhattacharya, S. S.; Lehmann, O. J.; Garway-Heath, D. F.; Hitchings, R. A.  
CS (1) Department of Ophthalmology, Hiroshima Univ Sch of Medicine, Hiroshima, Japan Japan  
SO ARVO Annual Meeting Abstract Search and Program Planner, (2002) Vol. 2002, pp. Abstract No. 3386. cd-rom.  
Meeting Info.: Annual Meeting of the Association For Research in Vision and Ophthalmology Fort Lauderdale, Florida, USA May 05-10, 2002  
DT Conference  
LA English

L5 ANSWER 14 OF 32 BIOSIS COPYRIGHT 2003 BIOLOGICAL ABSTRACTS INC.  
AN 2003:155032 BIOSIS  
DN PREV200300155032  
TI Investigating the Association Between OPA1 Polymorphisms and Glaucoma: Comparison Between Normal Tension Glaucoma and High Tension Primary Open Angle Glaucoma.  
AU Aung, T. (1); Ocaka, L.; Ebenezer, N. D.; Morris, A. G.; Brice, G.; Child, A. H.; Hitchings, R. A.; Lehmann, O. J.; Bhattacharya, S. S.

DT Conference  
LA English

L5 ANSWER 15 OF 32 MEDLINE  
AN 2001417813 MEDLINE  
DN 21334296 PubMed ID: 11440989  
TI Spectrum, frequency and penetrance of **OPA1** mutations in dominant optic atrophy.  
AU Toomes C; Marchbank N J; Mackey D A; Craig J E; Newbury-Ecob R A; Bennett C P; Vize C J; Desai S P; Black G C; Patel N; Teimory M; Markham A F; Inglehearn C F; Churchill A J  
CS Molecular Medicine Unit, Clinical Sciences Building, University of Leeds, St James's University Hospital, Leeds, LS9 7TF, UK.  
SO HUMAN MOLECULAR GENETICS, (2001 Jun 15) 10 (13) 1369-78.  
Journal code: 9208958. ISSN: 0964-6906.  
CY England: United Kingdom  
DT Journal; Article; (JOURNAL ARTICLE)  
LA English  
FS Priority Journals  
EM 200108  
ED Entered STN: 20010903  
Last Updated on STN: 20010903  
Entered Medline: 20010830

L5 ANSWER 16 OF 32 BIOSIS COPYRIGHT 2003 BIOLOGICAL ABSTRACTS INC.  
AN 2001:156289 BIOSIS  
DN PREV200100156289  
TI The pupil in dominant optic atrophy.  
AU Bremner, Fion D. (1); Tomlin, Elizabeth A.; Shallo-Hoffmann, Josephine; Votruba, Marcella; Smith, Stephen E.  
CS (1) Department of Neuroophthalmology, National Hospital for Neurology and Neurosurgery, Queen Square, London, WC1N 3BG: fdbremner@doctors.org.uk UK  
SO IOVS, (March, 2001) Vol. 42, No. 3, pp. 675-678. print.  
DT Article  
LA English  
SL English

L5 ANSWER 17 OF 32 BIOSIS COPYRIGHT 2003 BIOLOGICAL ABSTRACTS INC.  
AN 2001:351846 BIOSIS  
DN PREV200100351846  
TI Spatial and temporal retinal expression of **OPA1** involved in autosomal dominant optic atrophy.  
AU Pesch, U. E. A. (1); Fries, J. E.; Alexander, C. (1); Wheeler-Schilling, T. H.; Kohler, K.; Wissinger, B. (1)  
CS (1) Molecular Genetics Laboratory, University Eye Hospital, Tuebingen Germany  
SO IOVS, (March 15, 2001) Vol. 42, No. 4, pp. S654. print.  
Meeting Info.: Annual Meeting of the Association for Research in Vision and Ophthalmology Fort Lauderdale, Florida, USA April 29-May 04, 2001  
DT Conference  
LA English  
SL English

L5 ANSWER 18 OF 32 BIOSIS COPYRIGHT 2003 BIOLOGICAL ABSTRACTS INC.  
AN 2001:348217 BIOSIS  
DN PREV200100348217  
TI Mutation screening of **OPA1** in dominant optic atrophy.  
AU Murton, N. J. (1); Toomes, C. (1); Mackey, D. A.; Inglehearn, C. F. (1); Churchill, A. J. (1)  
CS (1) Molecular Medicine Unit, St James's University Hospital, Leeds UK  
SO IOVS, (March 15, 2001) Vol. 42, No. 4, pp. S650. print.  
Meeting Info.: Annual Meeting of the Association for Research in Vision and Ophthalmology Fort Lauderdale, Florida, USA April 29-May 04, 2001  
DT Conference

LN PREV200100343623  
TI A first locus for non syndromic autosomal recessive optic atrophy (OAR1).  
AU Kaplan, J. (1); Barber, F. (1); Gerber, S. (1); Rozet, J. M. (1);

CS Perrault, I. (1); Ghazi, I. (1); Ducrocq, D. (1); Hanein, S. (1); Lufier, J. L.; Munnich, A. (1)

SO (1) Genetics, INSERM U393, Hopital des Enfants Malades, Paris France

IOVS, (March 15, 2001) Vol. 42, No. 4, pp. S650. print.

Meeting Info.: Annual Meeting of the Association for Research in Vision and Ophthalmology Fort Lauderdale, Florida, USA April 29-May 04, 2001

DT Conference

LA English

SL English

L5 ANSWER 20 OF 32 BIOSIS COPYRIGHT 2003 BIOLOGICAL ABSTRACTS INC.

AN 2001:348218 BIOSIS

DN PREV200100348218

TI Nuclear gene **OPA1** encoding a mitochondrial dynamin-related protein is mutated in dominant optic atrophy.

AU Delettre, C. (1); Lenaers, G.; Griffoin, J. M. (1); Arnaud, B.; Dollfus, H.; Kaplan, J.; Lorenz, B.; Van de Kamp, J.; Belenguer, P.; Hamel, C. P. (1)

CS (1) Laboratoire de Neurobiologie de l'Audition, Inserm U. 254, Montpellier France

SO IOVS, (March 15, 2001) Vol. 42, No. 4, pp. S650. print.

Meeting Info.: Annual Meeting of the Association for Research in Vision and Ophthalmology Fort Lauderdale, Florida, USA April 29-May 04, 2001

DT Conference

LA English

SL English

L5 ANSWER 21 OF 32 MEDLINE DUPLICATE 6

AN 2002082743 MEDLINE

DN 21668183 PubMed ID: 11810270

TI Mutation spectrum and splicing variants in the **OPA1** gene.

AU Delettre C; Griffoin J M; Kaplan J; Dollfus H; Lorenz B; Faivre L; Lenaers G; Belenguer P; Hamel C P

CS Inserm U. 254, 71, rue de Navacelles, 34090 Montpellier, France.

SO HUMAN GENETICS, (2001 Dec) 109 (6) 584-91.

Journal code: 7613873. ISSN: 0340-6717.

CY Germany: Germany, Federal Republic of

DT Journal; Article; (JOURNAL ARTICLE)

LA English

FS Priority Journals

EM 200202

ED Entered STN: 20020128

Last Updated on STN: 20030105

Entered Medline: 20020215

L5 ANSWER 22 OF 32 BIOSIS COPYRIGHT 2003 BIOLOGICAL ABSTRACTS INC.

AN 2001:348053 BIOSIS

DN PREV200100348053

TI The gene responsible for autosomal dominant optic atrophy, **OPA1** is not associated with normal tension glaucoma.

AU Ocaka, L. (1); Aung, T. (1); Thiselton, D. L. (1); Francis, P. J. (1); Aragon-Martin, J. (1); Payne, A. (1); Child, A. H.; Votruba, M. (1); Hitchings, R. A.; Bhattacharya, S. S. (1)

CS (1) Molecular Genetics, Inst of Ophthalmology, London UK

SO IOVS, (March 15, 2001) Vol. 42, No. 4, pp. S564. print.

Meeting Info.: Annual Meeting of the Association for Research in Vision and Ophthalmology Fort Lauderdale, Florida, USA April 29-May 04, 2001

DT Conference

LA English

SL English

AU The gene responsible for dominant optic atrophy, **OPA1** is not associated with normal tension glaucoma. High prevalence of dominant optic atrophy in the Ashkenazi population provides evidence for a founder effect.

Thiselton D L; Alexander C; Morris A; Brooks S; Rosenberg T; Eiberg H; Kier B; Kier P; Bhattacharya S S; Votruba M

CS Department of Molecular Genetics, Institute of Ophthalmology, University  
College London, EC1 V 9EL, UK.. dthiselt@hgmp.mrc.ac.uk  
SO HUMAN GENETICS, (2001 Nov) 109 (5) 498-502.  
Journal code: 7613873. ISSN: 0340-6717.  
CY Germany: Germany, Federal Republic of  
DT Journal; Article; (JOURNAL ARTICLE)  
LA English  
FS Priority Journals  
EM 200112  
ED Entered STN: 20011206  
Last Updated on STN: 20030105  
Entered Medline: 20011220

LS ANSWER 24 OF 32 BIOSIS COPYRIGHT 2003 BIOLOGICAL ABSTRACTS INC.  
AN 2000:367518 BIOSIS  
DN PREV200000367518  
TI Characterisation of physically localised candidate transcripts at the  
locus for dominant optic atrophy (OPA1).  
AU Thiselton, D. L. (1); Alexander, C. (1); Votruba, M. (1); Pesch, U.;  
Bhattacharya, S. S. (1); Wissinger, B.  
CS (1) Molecular Genetics, Institute of Ophthalmology, UCL, London UK  
SO European Journal of Neuroscience, (2000) Vol. 12, No. Supplement 11, pp.  
230. print.  
Meeting Info.: Meeting of the Federation of European Neuroscience  
Societies Brighton, UK June 24-28, 2000  
ISSN: 0953-816X.

DT Conference  
LA English  
SL English

LS ANSWER 25 OF 32 MEDLINE DUPLICATE 8  
AN 2001019513 MEDLINE  
DN 20472324 PubMed ID: 11017080  
TI OPA1, encoding a dynamin-related GTPase, is mutated in autosomal  
dominant optic atrophy linked to chromosome 3q28.  
AU Alexander C; Votruba M; Pesch U E; Thiselton D L; Mayer S; Moore A;  
Rodriguez M; Kellner U; Leo-Kottler B; Auburger G; Bhattacharya S S;  
Wissinger B  
CS Department of Molecular Genetics, Institute of Ophthalmology, University  
College London, London, UK.  
SO NATURE GENETICS, (2000 Oct) 26 (2) 211-5.  
Journal code: 9216904. ISSN: 1061-4036.  
CY United States  
DT Journal; Article; (JOURNAL ARTICLE)  
LA English  
FS Priority Journals  
OS GENBANK-AB011139  
EM 200011  
ED Entered STN: 20010322  
Last Updated on STN: 20010322  
Entered Medline: 20001108

LS ANSWER 26 OF 32 MEDLINE DUPLICATE 9  
AN 2001019512 MEDLINE  
DN 20472323 PubMed ID: 11017079  
TI Nuclear gene OPA1, encoding a mitochondrial dynamin-related  
protein, is mutated in dominant optic atrophy.  
AU Delettre C; Lenaers G; Griffioen J M; Gigarel N; Lorenzo C; Belenguer P;  
Pelloquin L; Grosgeorge J; Turc-Carel C; Perret E; Astarie-Dequeker C;  
Lasquellec L; Arnaud B; Ducommun B; Kaplan J; Hamel C P  
CS Inserm U. 254, Laboratoire de Neurobiologie de l'audition, Montpellier,

LA English  
FS Priority Journals  
OS GENBANK AB006965; GENBANK AB011139; GENBANK AC023594; GENBANK AC025847;

EM GENBANK-J05620; GENBANK-L00007; GENBANK-Y07891; GENBANK-Z75  
200011  
ED Entered STN: 20010322  
Last Updated on STN: 20010322  
Entered Medline: 20001108

L5 ANSWER 27 OF 32 USPATFULL  
AN 1999:33783 USPATFULL  
TI Reciprocal subtraction differential display  
IN Fisher, Paul B., Scarsdale, NY, United States  
PA The Trustees of Columbia University in the City of New York, New York,  
NY, United States (U.S. corporation)  
PI US 5882874 19990316  
AI US 1998-32684 19980227  
DT Utility  
FS Granted  
LN.CNT 1353  
INCL INCLM: 435/006.000  
INCLS: 435/091.210  
NCL NCLM: 435/006.000  
NCLS: 435/091.210  
IC [6]  
ICM: C12Q001-68  
EXF 435/6; 435/172.3; 435/91.2; 435/91.21  
CAS INDEXING IS AVAILABLE FOR THIS PATENT.

L5 ANSWER 28 OF 32 PCTFULL COPYRIGHT 2003 Univentio  
AN 1999043844 PCTFULL ED 20020515  
TIEN RECIPROCAL SUBTRACTION DIFFERENTIAL DISPLAY  
TIFR IDENTIFICATION DIFFERENTIELLE A SOUSTRACTION RECIPROQUE  
IN FISHER, Paul, B.  
PA THE TRUSTEES OF COLUMBIA UNIVERSITY IN THE CITY OF NEW YORK;  
FISHER, Paul, B.  
LA English  
DT Patent  
PI WO 9943844 A1 19990902  
DS W: AU CA JP MX US AT BE CH CY DE DK ES FI FR GB GR IE IT LU MC  
NL PT SE  
AI WO 1999-US4323 A 19990226  
PRAI US 1998-09/032,684 19980227  
US 1998-09/185,115 19981103  
US 1998-09/197,889 19981123  
ICM C12P021-02  
ICS C12Q001-68; C12N015-11

L5 ANSWER 29 OF 32 MEDLINE DUPLICATE 10  
AN 1998455021 MEDLINE  
DN 98455021 PubMed ID: 9783700  
TI Clinical features, molecular genetics, and pathophysiology of dominant  
optic atrophy.  
AU Votruba M; Moore A T; Bhattacharya S S  
CS Department of Molecular Genetics, Institute of Ophthalmology, University  
College London, UK.  
SO JOURNAL OF MEDICAL GENETICS, (1998 Oct) 35 (10) 793-800. Ref: 96  
Journal code: 2985087R. ISSN: 0022-2593.  
CY ENGLAND: United Kingdom  
DT Journal; Article; (JOURNAL ARTICLE)  
General Review; (REVIEW)  
(REVIEW, TUTORIAL)  
LA English  
FS Priority Journals

L5 ANSWER 30 OF 32 MEDLINE  
AN 1998372640 MEDLINE  
DN 98372640 PubMed ID: 9708909

TI Isolation of a cDNA for a ~~tel~~ 120-kDa GTP-binding protein, expressed in  
motor **neurons** in the salmon brain.  
AU Kubokawa K; Miyashita T; Kubo Y  
CS Department of Molecular Biology, Ocean Research Institute, University of  
Tokyo, Nakano, Japan.  
SO FEBS LETTERS, (1998 Jul 17) 431 (2) 231-5.  
Journal code: 0155157. ISSN: 0014-5793.  
CY Netherlands  
DT Journal; Article; (JOURNAL ARTICLE)  
LA English  
FS Priority Journals  
OS GENBANK-AB012720  
EM 199809  
ED Entered STN: 19980917  
Last Updated on STN: 20000303  
Entered Medline: 19980908

L5 ANSWER 31 OF 32 MEDLINE DUPLICATE 11  
AN 1998141131 MEDLINE  
DN 98141131 PubMed ID: 9490303  
TI Demonstration of a founder effect and fine mapping of dominant optic  
atrophy locus on 3q28-qter by linkage disequilibrium method: a study of 38  
British Isles pedigrees.  
AU Votruba M; Moore A T; Bhattacharya S S  
CS Department of Molecular Genetics, Institute of Ophthalmology, London, UK..  
mvotruba@hgmp.mrc.ac.uk  
SO HUMAN GENETICS, (1998 Jan) 102 (1) 79-86.  
Journal code: 7613873. ISSN: 0340-6717.  
CY GERMANY: Germany, Federal Republic of  
DT Journal; Article; (JOURNAL ARTICLE)  
LA English  
FS Priority Journals  
EM 199803  
ED Entered STN: 19980312  
Last Updated on STN: 20000303  
Entered Medline: 19980303

L5 ANSWER 32 OF 32 BIOSIS COPYRIGHT 2003 BIOLOGICAL ABSTRACTS INC.  
AN 1998:111667 BIOSIS  
DN PREV199800111667  
TI Demonstration of a founder effect and linkage disequilibrium mapping in  
the refinement of the dominant optic atrophy locus (**OPA1**).  
AU Votruba, M. (1); Moore, A. T.; Bhattacharya, S. S.  
CS (1) Dep. Molecular Genetics, Inst. Ophthalmol., UCL, London UK  
SO American Journal of Human Genetics, (Oct., 1997) Vol. 61, No. 4 SUPPL.,  
pp. A299.  
Meeting Info.: 47th Annual Meeting of the American Society of Human  
Genetics Baltimore, Maryland, USA October 28-November 1, 1997  
ISSN: 0002-9297.  
DT Conference  
LA English

=> d 12 1-3

L2 ANSWER 1 OF 3 USPATFULL  
AN 2003:44371 USPATFULL  
TI Combined growth factor-deleted and thymidine kinase-deleted vaccinia  
virus vector  
IN McCart, J. Andrea, Toronto, CANADA  
Bartlett, David L., Pittsburgh, PA, UNITED STATES  
..

DT Utility;  
FS APPLICATION  
LN.CNT 2762

INCL INCLM: 424/186.100  
INCLS: 435/456.000; 435/235.100  
NCL NCLM: 424/186.100  
NCLS: 435/456.000; 435/235.100  
IC [7]  
ICM: A61K039-12  
ICS: C12N015-863; C12N007-00  
CAS INDEXING IS AVAILABLE FOR THIS PATENT.

L2 ANSWER 2 OF 3 USPATFULL  
AN 2003:3547 USPATFULL  
TI Trans-viral vector mediated gene transfer to the retina  
IN Wakefield, John, Birmingham, AL, UNITED STATES  
Bennett, Jean, Bryn Mawr, PA, UNITED STATES  
PA Tranzyme, Inc. (U.S. corporation)  
PI US 2003003582 A1 20030102  
AI US 2002-140227 A1 20020507 (10)  
PRAI US 2001-340116P 20011102 (60)  
US 2001-289459P 20010508 (60)  
DT Utility  
FS APPLICATION  
LN.CNT 2316  
INCL INCLM: 435/456.000  
INCLS: 424/093.200  
NCL NCLM: 435/456.000  
NCLS: 424/093.200  
IC [7]  
ICM: A61K048-00  
ICS: C12N015-867  
CAS INDEXING IS AVAILABLE FOR THIS PATENT.

L2 ANSWER 3 OF 3 PCTFULL COPYRIGHT 2003 Univentio  
AN 1996038552 PCTFULL ED 20020514  
TIEN POST-TRANSCRIPTIONAL GENE REGULATION BY SELENIUM  
TIFR REGULATION POST-TRANSCRIPTIONNELLE DE GENES PAR LE SELENIUM  
IN LEONARD, Jack, L.;  
NEWBURGER, Peter, E.  
PA UNIVERSITY OF MASSACHUSETTS MEDICAL CENTER  
LA English  
DT Patent  
PI WO 9638552 A1 19961205  
DS W: AU CA JP KR AT BE CH DE DK ES FI FR GB GR IE IT LU MC NL PT  
SE  
AI WO 1996-US7496 A 19960523  
PRAI US 1995-8/454,028 19950530  
ICM C12N015-09  
ICS C12N015-11; C12N015-63; C12P021-02